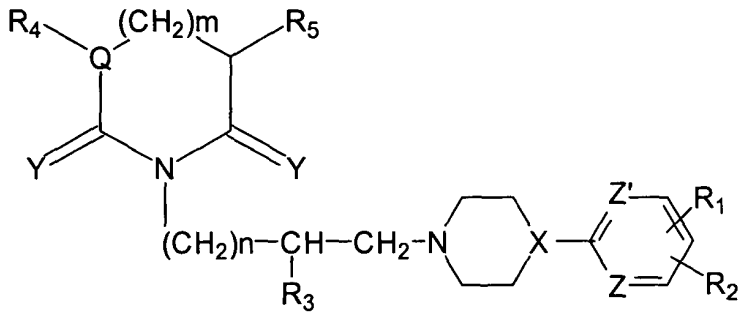
In the claims:

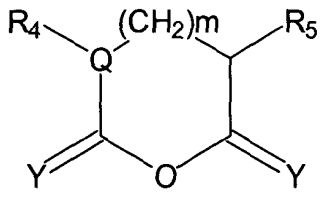
Kindly amend claims 44 and 45 to read as follows:

44. (Four Times Amended) A method for making a compound having the structure of Formula I



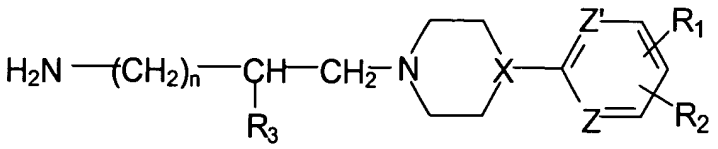
(I)

its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein **Y** is O or S; **Q, Z** and **Z'** are independently CH; X is CH or N; **m**=0; **n**=0-4; $\mathbf{R_1}$, $\mathbf{R_2}$ are independently selected from: F, Br, OCH_3, OC_2H_5, OCH_2CF_3, SCF_3, C_2H_5, CF_3, isopropyloxy, and cyclopropyl; and R_3, R_4 and R_5 are independently H, $C_{1\text{-}3}$ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI'



(VI')

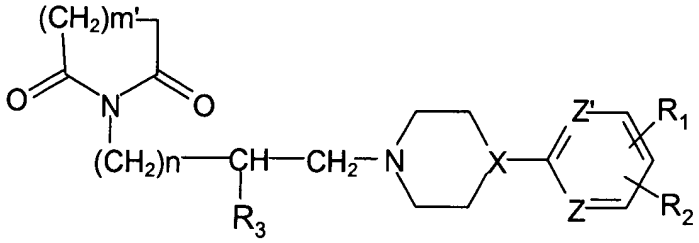
with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride



(V)

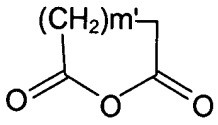
thereby to produce the compound of Formula I.

45. (Thrice Amended) A method for making a compound having the structure of Formula II



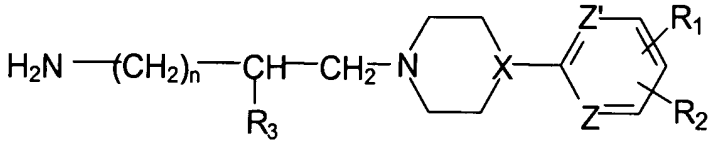
(II)

its pharmaceutically acceptable salts, enantiomers, diasteromers, or N-oxides, wherein X is CH or N; Z and Z' are independently CH; n= 0-4; m' = 1; R_1, R_2 are independently selected from: F, Br, OCH_3, OC_2H_5, OCH_2CF_3, SCF_3, isopropyloxy, and cyclopropyl; and R_3 is independently H, C_{1-3} alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI



(VI)

with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride

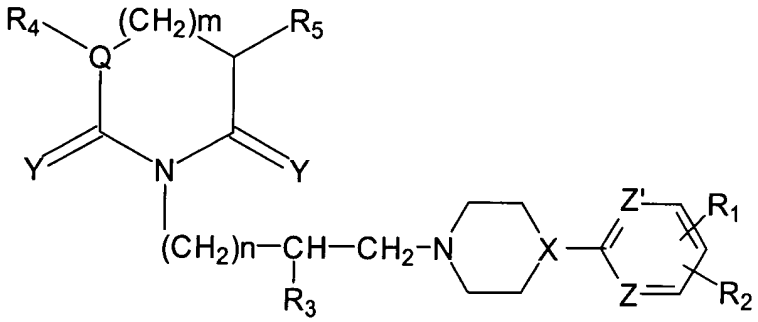


(V)

thereby to produce the compound of Formula II.

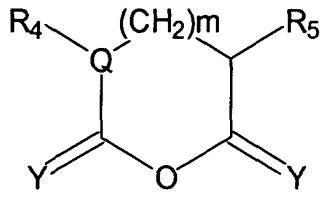
Please add new claims 48 and 49.

48. (New) A method for making a compound having the structure of Formula I



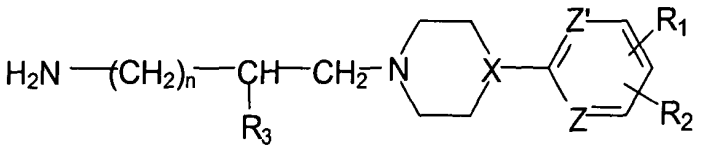
(I)

its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein **Y** is O or S; **Q, Z** and **Z′** are independently CH; X is CH or N; **m**=1-3; **n**=0-4; $\mathbf{R_1}$, $\mathbf{R_2}$ are independently selected from: F, Cl, Br, OC_2H_5, OCH_2CF_3, SCF_3, CH_3, C_2H_5, CF_3, isopropyloxy, and cyclopropyl; and R_3, R_4 and R_5 are independently H, $C_{1\text{-}3}$ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI′



(VI')

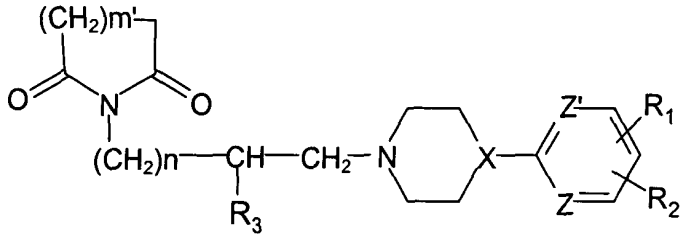
with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride



(V)

thereby to produce the compound of Formula I.

49. (New) A method for making a compound having the structure of Formula II



(II)

its pharmaceutically acceptable salts, enantiomers, diasteromers, or N-oxides, wherein X is CH or N; Z and Z' are independently CH; n= 0-4; m' = 2-4; R_1, R_2 are independently selected from: F, Cl, Br, OC_2H_5, OCH_2CF_3, SCF_3, CH_3, isopropyloxy, and cyclopropyl; and R_3 is independently H, C_{1-3} alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI